

in-part of USSN 07/619,131 filed November 27, 1990, which is a continuation-in-part of USSN 07/444,745 filed December 1, 1989.

New claim 98 is directed to a cDNA sequence encoding mature human lactoferrin as shown in Fig. 1 (SEQ. ID. No. 1). Support for the claim is provided by e.g., Fig. 1 and pp. 42-44 of the present application, and by e.g., Fig. 1 and pp. 25-27 of the earliest application from which priority is claimed, USSN 07/444,745 (filed December 1, 1989). By specifying that the claimed cDNA sequence comprises the region encoding the mature protein, it is meant that the region encoding the signal sequence need not be present, as explained in the present specification at p. 46, lines 21-23 and the '745 priority application at p. 26. lines 25-27.

New claim 98 is added to cause declaration of an interference between the present application and a US application corresponding to WO 90/13642, which claims priority from USSN 07/348,270 (filed May 5, 1989). WO 90/13642 (hereinafter the '642 PCT application) names Headon, Conneely and O'Malley as inventors and is assigned on its face to Granada Biosciences (although present assignment of a corresponding US application is unknown). A copy of the '642 PCT application is attached. It is believed that a Notice of Allowability has recently occurred in a US application corresponding to the '642 PCT application.

Although the present applicants have no way of knowing exactly what claims may have been allowed in a US application corresponding to the '642 PCT application, the '642 PCT application is clearly directed to closely related subject matter (i.e., cDNA sequences encoding human lactoferrin) to claim 98 in the present application. For example, claim 1 in the '642 PCT application reads,

A cDNA sequence coding for human lactoferrin protein, comprising the DNA sequence of Fig. 2.

The mature coding region of the DNA sequence in Fig. 2 of the '642 PCT application and that of Fig. 1 (SEQ. ID. No. 1) of the present application, respectively, encode mature human lactoferrin proteins of 692 and 691 amino acids. The respective

proteins are believed to differ at only three places<sup>1</sup> as follows.

aa#	Fig. 1 (present application)	Fig. 2 ('642 PCT application)
130	Threonine	Isoleucine
151	Glutamine	Alanine and Arginine
403	Cysteine	Glycine

Under the provisions of 37 CFR §1.604, applicants seek to have an interference declared with a US application corresponding to the '642 PCT application. Applicants present the following proposed count 1, which is the same as new claim 98 in the present application:

An isolated cDNA sequence comprising the region of the cDNA sequence of Fig. 1 (SEQ. ID. No. 1) encoding mature human lactoferrin protein.

According to 37 CFR §1.601(i), an interference may be declared between two or more pending applications naming different inventors when, in the opinion of the Examiner, the applications contain claims for the same patentable invention as defined in §1.601(n). Here, declaration of an interference is particularly appropriate because the respective priority dates of the two applications at issue are only about six months apart. Although applicants are unable to identify a claim from a US application corresponding to the '642 PCT application, which corresponds to the count (due to pending US applications being maintained in confidence), they request the Examiner to review

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<sup>1</sup>Note that amino acid position numbers are those of Fig. 1 in the present application, in which the first amino acid of the mature protein is designated amino acid 1. The sequence in Fig. 2 of the '642 PCT application is numbered by a different convention in which the first amino acid of the signal peptide is designated 1.

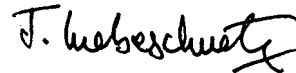
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the allowed claims in the US application corresponding to the '642 PCT application to determine whether any such claim corresponds to the proposed count.

If any such claim in a US application corresponding to the '642 PCT application does correspond to the proposed count, the Examiner is requested to declare an interference and stay issuance of the US application corresponding to the '642 PCT application.

Respectfully submitted,



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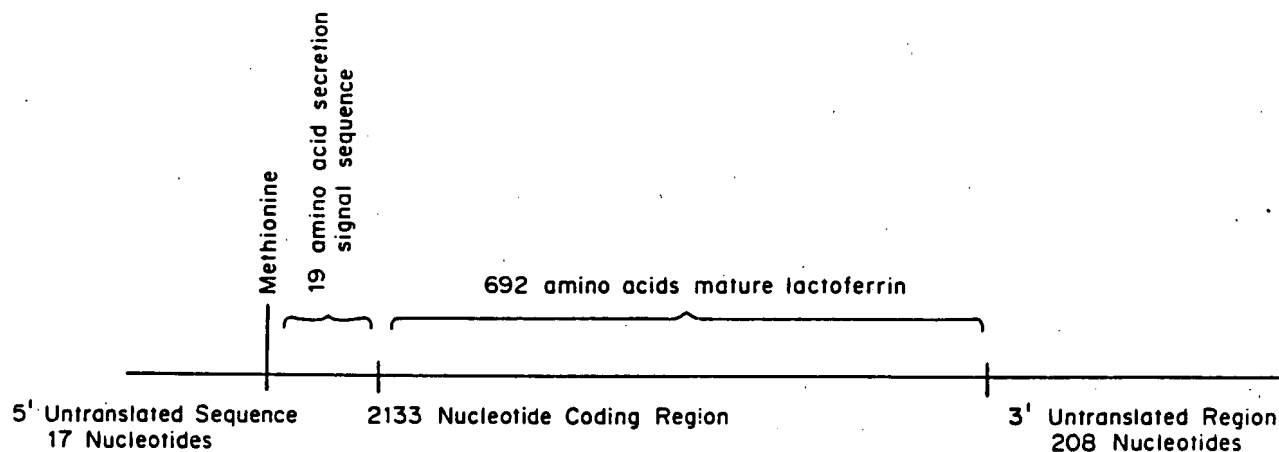
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## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 5 : <b>C12N 15/12</b>	<b>A1</b>	(11) International Publication Number: <b>WO 90/13642</b> (43) International Publication Date: 15 November 1990 (15.11.90)
<p>(21) International Application Number: PCT/US90/02356</p> <p>(22) International Filing Date: 27 April 1990 (27.04.90)</p> <p>(30) Priority data: 348,270      5 May 1989 (05.05.89)      US</p> <p>(71) Applicant: GRANADA BIOSCIENCES, INC. [US/US]; 10900 Richmond Avenue, Houston, TX 77242 (US).</p> <p>(72) Inventors: HEADON, Denis, R. ; Lishin, Agnacuerra, Dangan Lower, Galway (IE). CONNEELY, Orma, M. ; 2601 South Braeswood, #1307, Houston, TX 77025 (US). O'MALLEY, Bert, W. ; 639 Ramblewood, Houston, TX 77079 (US).</p> <p>(74) Agent: BOULWARE, Margaret, A.; One Riverway, Suite 1100, Houston, TX 77056-1903 (US).</p>		<p>(81) Designated States: AT (European patent), AU, BE (Euro- + pean patent), CA, CH (European patent), DE (Euro- pean patent), DK (European patent), ES (European pat- ent), FR (European patent), GB (European patent), IT (European patent), JP, LU (European patent), NL (Eu- ropean patent), SE (European patent).</p> <p><b>Published</b> <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i></p> <p>CUSTOMER COPY Complies with Copyright Law CAS Document Delivery Service 1-614-447-3670 <b>74-064532</b> <b>90486892M</b></p>

(54) Title: HUMAN LACTOFERRIN cDNA SEQUENCE

## Scheme of Human Lactoferrin (hLF) cDNA



Complete Human Lactoferrin : 711 amino acids  
Mature Human Lactoferrin : 692 amino acids

## (57) Abstract

The verified cDNA sequence for human lactoferrin (hLF) protein has been discovered. The cDNA sequence can be used to prepare recombinant human lactoferrin for therapeutic and nutritional applications. Regions of the cDNA such as the Fe binding sites can be used to make an hLF polypeptide product.

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## HUMAN LACTOFERRIN cDNA SEQUENCE

BACKGROUND OF THE INVENTION

Lactoferrin is an iron-binding glycoprotein found in milk and other secretions and body fluids. It is one of a number of iron binding proteins, sometimes referred to as transferrins, and is involved in iron binding and delivery in mammals.

Lactoferrin has been implicated as a factor in resistance against enteritis infections in suckled newborns. These bacteriocidal/bacteriostatic actions are considered to be due at least in part to the iron binding properties of lactoferrin. Lactoferrin decreases the iron availability to iron requiring microorganisms and thereby interferes with their growth and reproduction. Lactoferrin is also considered to have antiviral properties and to have other potential therapeutic applications.

Human lactoferrin (hLF) has a high affinity for iron and two  $\text{Fe}^{3+}$  cations can be found per molecule. The complete hLF protein has been subjected to amino acid sequencing and is reported to have 703 amino acids. There are two glycosylation sites. Metz-Boutigue et al; "Human Lactotransferrin: Amino Acid Sequence and Structural Comparisons With Other Transferrins", Eur. J. Biochem., vol. 145, pp. 659-676 (1984). Anderson et al; "Structure of Human Lactoferrin at 3.2 - A Resolution", Proc. Nat'l. Acad. Sci. USA, vol. 84, pp. 1769-1773 (April 1987).

In other studies, a cloned cDNA probe for amino acids 428 to 703 of the lactoferrin protein was isolated. The cDNA sequence was in general agreement with the earlier analysis of the amino acid sequence of the protein. Rado, et al; "Isolation of Lactoferrin cDNA From a Human Myeloid

Library and Expression of mRNA During Normal and Leukemic Myelopoiesis", Blood, vol. 79, no. 4, pp. 989-993 (Oct. 1987). The probe was reported to encompass approximately 40% of the coding region and the 3' terminus. The full confirmed cDNA sequence has not been reported.

#### SUMMARY OF THE INVENTION

The invention is the verified cDNA sequence for human lactoferrin which includes an open reading frame of 2133 nucleotides coding for a protein of 711 amino acids. These 711 amino acids include 19 amino acids corresponding to a secretion signal peptide sequence followed by 692 amino acids of mature human lactoferrin. The cDNA sequence and deduced amino acid sequence differ from the previously published data. The confirmation of the cDNA sequence and the deduced amino acid have been proven by multiple confirmation procedures.

These are:

1. Multiple sequence analyses.
2. Comparison of the amino acid sequence deduced from the cDNA with that of human LF generated by conventional amino acid sequencing of hLF isolated from milk. The unique cDNA sequence which encodes the human lactoferrin protein has a variety of applications as known and indicated in the literature.
3. Transcription and translation of hLF protein from the cDNA with positive identification using an anti hLF antibody.

The present invention of the cDNA sequence can be used to prepare recombinant human lactoferrin, thus making available a source of human protein for therapeutic and nutritional applications. The confirmed cDNA of this invention can be used in an appropriate cloning vehicle to replicate the cDNA sequence. Also, the cDNA can be incorporated into a vector system for human lactoferrin

expression. This invention is not limited to any particular uses of the cDNA. The recombinant hLF being a human protein derived by recombinant techniques can be used in a variety of applications. The gene can be transferred to mammalian systems such as cows and other agriculturally important animals and expressed in milk. The incorporation of a human lactoferrin and expression in the milk of animals can combat an iron deficiency typical in piglets. The inclusion of the human lactoferrin gene with expression should improve an animal's disease resistance to bacterial and viral infection. The tissue specific expression of human lactoferrin in mammary glands, for instance, would impart the bacteriocidal and viracidal benefit of the expressed gene to young feeding on the milk and would provide a production means for the secreted protein for therapeutic use.

The gene can be placed in the appropriate cloning vector for the production of hLF. The hLF produced by recombinant methods can be used in a variety of products including human or animal foods, as therapeutic additives to enhance iron transport and delivery, and for the viracidal and bacteriocidal qualities, as additives for eye-drops, contact lens and other eye care solutions, topical skin care products, ear drops, mouthwashes, chewing gum and toothpaste. The recombinant hLF would provide a safe, naturally occurring product which can be topically applied as well as ingested safely.

#### BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 is a schematic drawing of the hLF cDNA including the locations of the 5' untranslated region, the secretion peptide signal sequence, mature lactoferrin and 3' untranslated region.

Fig. 2 is the cDNA sequence with deduced amino acids for the human lactoferrin protein and signal peptide.



sequence.

Fig. 3 is a schematic representation of an autoradiograph of recombinant human lactoferrin protein expressed from the complete cDNA.

Fig. 4 is a schematic representation of an autoradiograph of the results of in vitro translation of a 2,140 bp human lactoferrin sequence and hLF protein in reticulocyte lysates.

#### DETAILED DESCRIPTION OF THE INVENTION

The full length cDNA encoding human lactoferrin (hLF) has been isolated, and the analysis has been completed. The cDNA sequence has been confirmed as human lactoferrin cDNA by comparison of the deduced amino acid sequence with the published amino acid sequence of human LF. The expression of lactoferrin was observed in an eucaryotic expression system from the cDNA and a plasmid vector. The presence of lactoferrin was confirmed by standard Western immunoblot analysis using anti-human lactoferrin antibodies and relative molecular mass measurement.

Fig. 1 is a schematic of the lactoferrin cDNA. The sequence can generally be described as an initial 5' untranslated region, 17 nucleotides in length. The next portion is 57 nucleotides which codes for the 19 amino acid secretion signal peptide starting with methionine. The next sequence of the cDNA codes for the mature human lactoferrin protein of 692 amino acids followed by the 3' untranslated region of 208 nucleotides which ends the cDNA. The complete sequence is 2,358 nucleotides in length. The hLF protein contains glycosylation sites. The hLF protein with secretion signal sequence has an expected molecular mass of 78,403 daltons and the mature hLF is 76,386 daltons without added carbohydrate from glycosylation.

Fig. 2 is the cDNA sequence with the deduced amino acids for the secretion signal peptide and the mature human lactoferrin protein. The numbers on Fig. 2 correspond to the nucleotides starting at the 5' end. There are binding sites for two iron atoms with four amino acids participating in the binding of each iron. The amino acids at positions Asp80, Tyr112, Tyr209, and His273 are required for coordination with one iron, and amino acids at positions Asp415, Tyr455, Tyr548, and His617 bind the other. There are two glycosylation sites at positions Asn157 and Asn498. The numbers refer to the deduced amino acid sequence. There are 25 amino acids per line of protein sequence (starting at nucleotide 18).

The nucleotide sequence analysis was performed on cDNA isolated from a human prostate cDNA library. The prostate cDNA library yielded a 2,140 bp cDNA which contained the complete 5' end including the untranslated portion and the signal sequence. The 3' end including the three amino acids at the carboxy terminal and the untranslated region were obtained as a 208 bp cDNA from both a monocyte cDNA library and human prostate cDNA library.

The data in Fig. 2 displays the full length cDNA sequence of this invention. The complete sequence including the 5' untranslated region and signal peptide have not been reported. Further, the previously reported amino acid sequence varies from the deduced amino acid sequence sequence for hLF of this invention. The following TABLE 1 is a summary of the differences of the amino acid sequence of the present invention and those reported by Metz-Boutigue et al, Eur. J. Biochem., vol. 45, pp. 659-6 (1984). For the purpose of this table, the numbering of the amino acids will be initiated with methionine at the start of the signal peptide sequence as amino acid #1.

TABLE 1  
COMPARISON OF AMINO ACID SEQUENCES  
HUMAN LACTOFERRIN

<u>Amino Acid Deduced from cDNA of hLF</u>	<u>Change</u>	<u>Metz-Boutique Sequence</u>
# 30 Thr	Substitution	Ala
# 48 Arg	Substitution	Lys
# 141 Arg	Insertion	NONE
# 155 Phe	Substitution	Leu
# 156 Leu	Substitution	Phe
# 170 Ala	Insertion	NONE
# 204 Ser	Substitution	Leu
# 206 Gln	Substitution	Lys
# 209 Tyr	Substitution	Lys
# 321 Lys	Substitution	Phe
# 386 Glu	Substitution	Gln
# 392 Ser	Substitution	Trp
# 410 Asp	Substitution	Asn
# 411-412	Deletion	13 Amino acids in protein sequence not in deduced amino acid sequence from cDNA.
# 532 Gln	Substitution	Glu
# 695 Lys	Substitution	Arg

Fig. 3 is the expression of human lactoferrin protein from the complete hLF cDNA. The complete ~~2358~~ bp hLF cDNA was ligated to the eucaryotic expression vector, p91023(B) at the EcoRI site downstream from the adenovirus major late promoter. This plasmid vector was provided by Genetics Institute (Cambridge, Massachusetts) and has been described in previous publications (Wong et al., Science 288, 810-815 (1985)). The hLF cDNA expression vector was transferred into COSM-6 monkey kidney cells using standard tissue culture transfection conditions (Wigler et al. (1979) "Transformation of Mammalian Cells with Genes from Procarotes and Eucaryotes"; Cell, 16:777-785). These COS cells do not normally express lactoferrin. Forty-eight hours after transfection, the cells were harvested and crude cell extracts were prepared. Positive identification of the human lactoferrin was made by standard Western immunoblot analysis of the proteins expressed in the cell extracts, as

well as those secreted into the cell growth medium using a commercially available antibody directed against human lactoferrin (Sigma). Proteins which bound to the anti-lactoferrin antibody were detected using radio-iodine labelled Protein A which reacts with the antibody. The immunoblots were autoradiographed to identify the human lactoferrin protein. Fig. 3 is an autoradiographic film showing the human lactoferrin expressed in four cell extracts prepared from tissue culture cells which were transfected with the lactoferrin cDNA expression vector (lanes 5 to 8). Lanes 5 to 8 show that the transfected cells all contain human lactoferrin (marked with an arrow) which is immunoreactive with the antilactoferrin antibody and is the same molecular weight as human lactoferrin ( $M_r = 78,403$  daltons). The control cells which were not transfected with the cDNA did not contain lactoferrin (lanes 3 and 4). Analysis of the growth medium showed that human lactoferrin was also secreted into the medium from transfected cells (lane 2) but not from control cells (lane 1).

In conclusion, the cDNA encodes a recombinant human lactoferrin protein which is similar to human lactoferrin protein isolated from milk as determined by molecular size comparisons and immunoreactivity with antihuman lactoferrin. Furthermore, the secretion signal peptide sequence is functional since the human lactoferrin is secreted into the growth medium of tissue culture cells which express the cDNA.

Fig. 4 is a schematic representation of the human lactoferrin protein precipitated after in vitro transcription and translation of the human lactoferrin cDNA. The 2140 bp cDNA was from the human prostate cDNA library and included the 5' untranslated region and the rest of the base pairs correlative to the cDNA sequence of Fig. 2 omitting the last 218 bp at the 3' terminus. The 2140 bp cDNA was ligated to the EcoRI site of the plasmid vector pGEM<sub>4</sub> (commercially available from Promega Biotech., Madison, WI 53711-5305) downstream from the SP<sub>6</sub> promoter. The plasmid

construct was linearized at the 3' end of the hLF cDNA using the restriction enzyme Hinc II or XBAI. The linear DNA template was then transcribed in vitro using purified SP<sub>6</sub> RNA polymerase in the presence of ribonucleotides as described in the manufacturers protocol (Promega Corporation 1988/1989 Catalogue and Applications Guide). The resultant mRNA was translated using 100ng mRNA template and micrococcal nuclease treated rabbit reticulocyte lysate (as described by Promega) in the presence of 75uCi <sup>35</sup>S methionine (800 ci/mmol, Amersham). In vitro synthesized lactoferrin was immunoprecipitated by incubating 100ul aliquots of translation reaction with 10ug of rabbit anti-human lactoferrin IgG (Sigma Chemical Company, St. Louis, MO 63178) for 2 hours at 4°C in 50mM Tris, pH7.5/0.15M NaCl/0.05% Tween-20 (1P-buffer). The reaction volume was 200ul. Immunoreactive lactoferrin was precipitated after incubation for 1 hour with 50ug of Protein A sepharose (Pharmacia, Upsalla, Sweden). Immunoprecipitation was carried out by centrifugation for 5 minutes at 10,000g and the precipitate was washed 5 times with 4 volumes of 1P buffer. Total translation products and immunoprecipitates were then subjected to electrophoresis in denaturing 7.5% polyacrylamide gels. After fixing in 50% Methanol, the gels were incubated in En<sup>3</sup>Hance (NEN, DuPont, Wilmington, DE 19801) for 1 hour and washed with distilled H<sub>2</sub>O. The gel was then dried under vacuum and exposed to Kodak X-OMAT XAR film at -70°C.

Lane 1 shows <sup>14</sup>C protein molecular weight markers used to estimate the size of the translated proteins. Lane 2 is a negative control which shows that no <sup>35</sup>S labelled proteins are translated in this system when no mRNA is added to the translation mix. Lanes 3 and 4 show the total translation products obtained when lactoferrin mRNA is added after preparation from two separate DNA templates. The major protein band (marked with an arrow) is human lactoferrin. This is the only band detected when the translation products are immunoprecipitated with anti human lactoferrin before

applying the protein to the gel (lane 6). The measurement of molecular mass by SDS-PAGE does not correspond to exact molecular weight due to secondary protein structure. However, the values are shifted in a correlative manner in comparison to the control. Analysis of the size of the translated lactoferrin is shown in Fig. 4. The protein migrated at the expected molecular mass of human lactoferrin (about 78Kd). The major bands in lanes 3 and 4 which migrate higher than the 68Kd marker band in the control lane correspond to expected molecular mass of hLF protein on SDS-PAGE.

In addition to using the entire cDNA sequence and deduced amino acids sequence, a polypeptide of less than the entire protein can be of value. For instance, the region between amino acids 74-275 contains an iron binding domain which may be used without the rest of the protein for biologically available iron or the bacteriocidal/viracidal qualities.

The cDNA sequence has been confirmed by corroboration of the sequence from three sources. The hLF cDNA was shown to encode lactoferrin by expression of the cDNA in a eucaryotic expression system and detection of the expressed lactoferrin protein by Western immunoblot analysis using specific lactoferrin antibodies.

1. A cDNA sequence coding for human lactoferrin protein, comprising the DNA sequence of Fig. 2.
2. A cDNA sequence coding for the functional equivalent of human lactoferrin comprising analogues of the cDNA sequence of Fig. 2.
3. A synthetic human lactoferrin comprising a product produced from the cDNA sequence of Fig. 2.
4. A synthetic functional equivalent of human lactoferrin comprising a product produced from analogues of the cDNA sequence of Fig. 2.
5. A method to produce synthetic human lactoferrin product comprising utilizing the cDNA sequence of Fig. 2.
6. A method to produce a synthetic functional equivalent of human lactoferrin comprising utilizing analogues of the cDNA sequence of Fig. 2.
7. A method to produce synthetic human lactoferrin and functional equivalents thereof of claims 5 and 6 utilizing a eucaryotic expression system.
8. A cDNA sequence comprising a portion of the cDNA of Fig. 2 coding for human lactoferrin protein including at least one of the iron binding domains with an Fe binding site.
9. A cDNA sequence of claim 8 wherein an analogue of a portion of the cDNA sequence codes for a functional equivalent of at least one iron binding domain with an Fe binding site.

10. A method comprising producing a portion of synthetic human lactoferrin of claim 8.

11. A method comprising producing a portion of synthetic functional equivalent of human lactoferrin of claim 9.

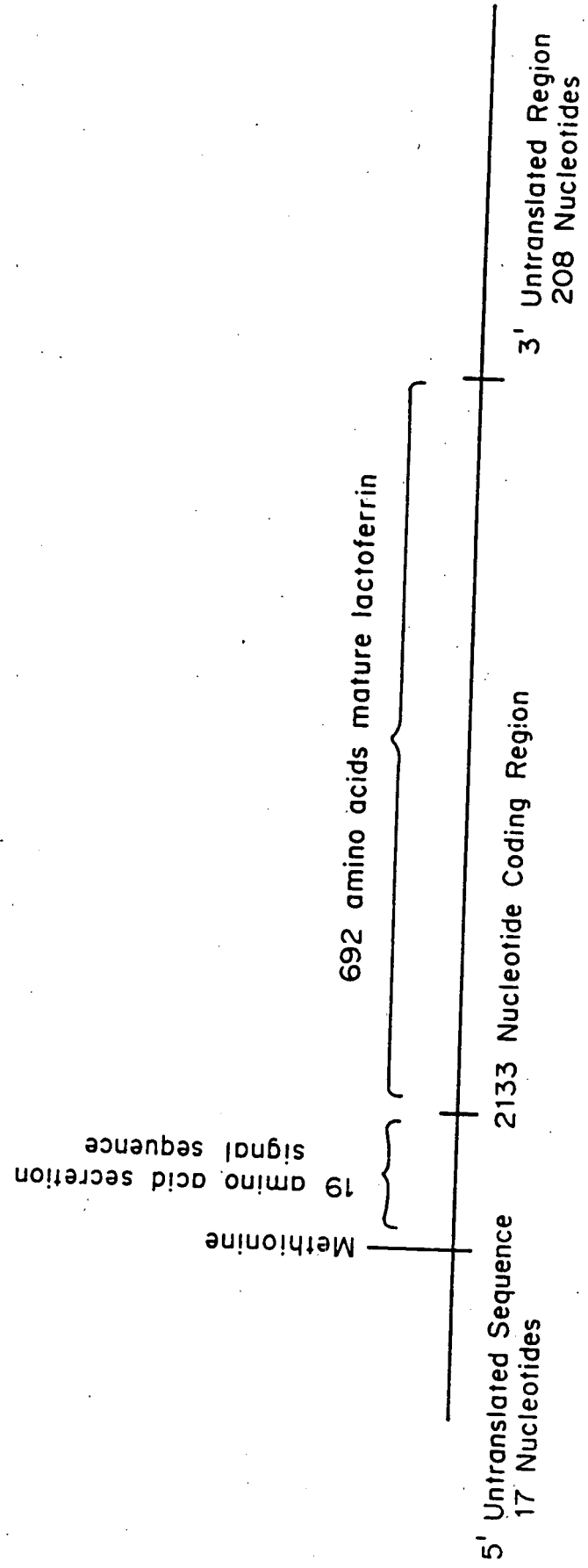
12. A synthetic human lactoferrin product comprising a portion of the synthetic human lactoferrin product produced from the cDNA sequence of Fig. 2 including at least one of the iron binding domains with an Fe binding site.

13. A synthetic human lactoferrin product comprising a synthetic functional equivalent of a portion of the human lactoferrin protein produced by an analogue of the cDNA sequence of Fig. 2 including at least one functional equivalent of the iron binding domains with a Fe binding site.



FIGURE 1

Scheme of Human Lactoferrin (hLF) cDNA



Complete Human Lactoferrin : 711 amino acids  
Mature Human Lactoferrin : 692 amino acids

2/7

**fig. 2**

Nice!

CDNA SEQUENCE AND AMINO ACID SEQUENCE  
OF HUMAN LACTOFERRIN

18  
 (2) (ATG AAA CTT GTC TTC CTC GTC CTG CTG TTC CTC GGG GCC CTC GGA CTG  
 met lys leu val phe leu val leu leu phe leu gly ala leu gly leu  
 1  
 66  
 TGT CTG GCT GGC CGT AGG AGA AGG AGT GTT CAG TGG TGC ACC GTA TCC  
 cys leu ala gly arg arg arg arg ser val gln trp cys thr val ser  
 17  
 114  
 CAA CCC GAG GCC ACA AAA TGC TTC CAA TGG CAA AGG AAT ATG AGA AGA  
 gln pro glu ala thr lys cys phe gln trp gln arg asn met arg arg  
 33  
 162  
 GTG CGT GGC CCT CCT GTC AGC TGC ATA AAG AGA GAC TCC CCC ATC CAG  
 val arg gly pro pro val ser cys ile lys arg asp ser pro ile gln  
 49  
 210  
 TGT ATC CAG GCC ATT GCG GAA AAC AGG GCC GAT GCT GTG ACC CTT GAT  
 cys ile gln ala ile ala glu asn arg ala asp ala val thr leu asp  
 65  
 258  
 GGT GGT TTC ATA TAC GAG GCA GGC CTG GCC CCC TAC AAA CTG CGA CCT  
 gly gly phe ile tyr glu ala gly leu ala pro tyr lys leu arg pro  
 81  
 306  
 GTA GCG GCG GAA GTC TAC GGG ACC GAA AGA CAG CCA CGA ACT CAC TAT  
 val ala ala glu val tyr gly thr glu arg gln pro arg thr his tyr  
 97  
 354  
 TAT GCC GTG GCT GTG GTG AAG AAG GGC GGC AGC TTT CAG CTG AAC GAA  
 tyr ala val ala val val lys lys gly gly ser phe gln leu asn glu  
 113  
 402  
 CTG CAA GGT CTG AAG TCC TGC CAC ACA GGC CTT CGC AGG ACC GCT GGA  
 leu gln gly leu lys ser cys his thr gly leu arg arg thr ala gly  
 129  
 450  
 TGG AAT GTG CCT ATA GGG ACA CTT CGT CCA TTC TTG AAT TGG ACG GGT  
 trp asn val pro ile gly thr leu arg pro phe leu asn trp thr gly  
 145

498  
CCA CCT GAG CCC ATT GAG GCA GCT GTG GCC AGG TTC TTC TCA GCC AGC  
pro pro glu pro ile glu ala ala val ala arg phe phe ser ala ser  
161

546  
TGT GTT CCC GGT GCA GAT AAA GGA CAG TTC CCC AAC CTG TGT CGC CTG  
cys val pro gly ala asp lys gly gln phe pro asn leu cys arg leu  
177

594  
TGT GCG GGG ACA GGG GAA AAC AAA TGT GCC TTC TCC TCC CAG GAA CCG  
cys ala gly thr gly glu asn lys cys ala phe ser ser gln glu pro  
193

642  
TAC TTC AGC TAC TCT GGT GCC TTC AAG TGT CTG AGA GAC GGG GCT GGA  
tyr phe ser tyr ser gly ala phe lys cys leu arg asp gly ala gly  
209

690  
GAC GTG GCT TTT ATC AGA GAG AGC ACA GTG TTT GAG GAC CTG TCA GAC  
asp val ala phe ile arg glu ser thr val phe glu asp leu ser asp  
225

738  
GAG GCT GAA AGG GAC GAG TAT GAG TTA CTC TGC CCA GAC AAC ACT CGG  
glu ala glu arg asp glu tyr glu leu leu cys pro asp asn thr arg  
241

786  
AAG CCA GTG GAC AAG TTC AAA GAC TGC CAT CTG GCC CGG GTC CCT TCT  
lys pro val asp lys phe lys asp cys his leu ala arg val pro ser  
257

834  
CAT GCC GTT GTG GCA CGA AGT GTG AAT GGC AAG GAG GAT GCC ATC TGG  
his ala val val ala arg ser val asn gly lys glu asp ala ile trp  
273

882  
AAT CTT CTC CGC CAG GCA CAG GAA AAG TTT GGA AAG GAC AAG TCA CCG  
asn leu leu arg gln ala gln gln lys phe gly lys asp lys ser pro  
289

930  
AAA TTC CAG CTC TTT GGC TCC CCT AGT GGG CAG AAA GAT CTG CTG TTC  
lys phe gln leu phe gly ser pro ser gly gln lys asp leu leu phe  
305

978  
AAG GAC TCT GCC ATT GGG TTT TCG AGG GTG CCC CCG AGG ATA GAT TCT  
lys asp ser ala ile gly phe ser arg val pro pro arg ile asp ser  
321

1026  
GGG CTG TAC CTT GGC TCC GGC TAC TTC ACT GCC ATC CAG AAC TTG AGG  
gly leu tyr leu gly ser gly tyr phe thr ala ile gln asn leu arg  
337

1074  
AAA AGT GAG GAG GAA GTG GCT GCC CGG CGT GCG CGG GTC GTG TGG TGT  
lys ser glu glu glu val ala ala arg arg ala arg val val trp cys  
353

1122  
GCG GTG GGC GAG CAG GAG CTG CGC AAG TGT AAC CAG TGG AGT GGC TTG  
ala val gly glu gln glu leu arg lys cys asn gln trp ser gly leu  
369

1170  
AGC GAA GGC AGC GTG ACC TGC TCC TCG GCC TCC ACC ACA GAG GAC TGC  
ser glu gly ser val thr cys ser ser ala ser thr thr glu asp cys  
385

1218  
ATC GCC CTG GTG CTG AAA GGA GAA GCT GAT GCC ATG AGT TTG GAT GGA  
ile ala leu val leu lys gly glu ala asp ala met ser leu asp gly  
401

1266  
GGA TAT GTG TAC ACT GCA GGC AAA TGT GGT TTG GTG CCT GTC CTG GCA  
gly tyr val tyr thr ala gly lys cys gly leu val pro val leu ala  
417

1314  
GAG AAC TAC AAA TCC CAA CAA AGC AGT GAC CCT GAT CCT AAC TGT GTG  
glu asn tyr lys ser gln gln ser ser asp pro asp pro asn cys val  
433

1362  
GAT AGA CCT GTG GAA GGA TAT CTT GCT GTG GCG GTG GTT AGG AGA TCA  
asp arg pro val glu gly tyr leu ala val ala val val arg arg ser  
449

1410  
GAC ACT AGC CTT ACC TGG AAC TCT GTG AAA GGC AAG AAG TCC TGC CAC  
asp thr ser leu thr trp asn ser val lys gly lys lys ser cys his  
465

1458  
ACC GCC GTG GAC AGG ACT GCA GGC TGG AAT ATC CCC ATG GGC CTG CTC  
thr ala val asp arg thr ala gly trp asn ile pro met gly leu leu  
481

1506  
TTC AAC CAG ACG GGC TCC TGC AAA TTT GAT GAA TAT TTC AGT CAA AGC  
phe asn gln thr gly ser cys lys phe asp glu tyr phe ser gln ser  
497

1554  
TGT GCC CCT GGG TCT GAC CCG AGA TCT AAT CTC TGT GCT CTG TGT ATT  
cys ala pro gly ser asp pro arg ser asn leu cys ala leu cys ile  
513

1602  
GGC GAC GAG CAG GCT GAG AAT AAG TGC GTG CCC AAC AGC AAT GAG AGA  
gly asp glu gln gly glu asn lys cys val pro asn ser asn glu arg  
529

5/7

1650

TAC TAC GGC TAC ACT GGG GCT TTC CGG TGC CTG GCT GAG AAT GCT GGA  
 tyr tyr gly tyr thr gly ala phe arg cys leu ala glu asn ala gly  
 545

1698

GAC GTT GCA TTT GTG AAA GAT GTC ACT GTC TTG CAG AAC ACT GAT GGA  
 asp val ala phe val lys asp val thr val leu gln asn thr asp gly  
 561

1746

AAT AAC AAT GAG GCA TGG GCT AAG GAT TTG AAG CTG GCA GAC TTT GCG  
 asn asn asn glu ala trp ala lys asp leu lys leu ala asp phe ala  
 577

1794

CTG CTG TGC CTC GAT GGC AAA CGG AAG CCT GTG ACT GAG GCT AGA AGC  
 leu leu cys leu asp gly lys arg lys pro val thr glu ala arg ser  
 593

1842

TTC CAT CTT GCC ATG GCC CCG AAT CAT GCC GTG GTG TCT CGG ATG GAT  
 cys his leu ala met ala pro asn his ala val val ser arg met asp  
 609

1890

AAG GTG GAA CGC CTG AAA CAG GTG CTG CTC CAC CAA CAG GCT AAA TTT  
 lys val glu arg leu lys gln val leu leu his gln gln ala lys phe  
 625

1938

GGG AGA AAT GGA TCT GAC TGC CCG GAC AAG TTT TGC TTA TTC CAG TCT  
 gly arg asn gly ser asp cys pro asp lys phe cys leu phe gln ser  
 641

1986

GAA ACC AAA AAC CTT CTG TTC AAT GAC AAC ACT GAG TGT CTG GCC AGA  
 glu thr lys asn leu leu phe asn asp asn thr glu cys leu ala arg  
 657

2034

CTC CAT GGC AAA ACA ACA TAT GAA AAA TAT TTG GGA CCA CAG TAT GTC  
 leu his gly lys thr thr tyr glu lys tyr leu gly pro gln tyr val  
 673

2082

GCA GGC ATT ACT AAT CTG AAA AAG TGC TCA ACC TCC CCC CTC CTG GAA  
 ala gly ile thr asn leu lys lys cys ser thr ser pro leu leu glu  
 689

2130

GCC TGT GAA TTC CTC AGG AAG TAA ) 2  
 ala cys glu phe leu arg lys ) \*\*\* ACCGAA GAAGATGGCC CAGCTCCCCA  
 705 ) 3 ) 4

2180

AGAAAGCCTC AGCCATTAC TGCCCCCAGC TCTTCTCCCC AGGTGTGTTG GGGCCTTGGC

2240

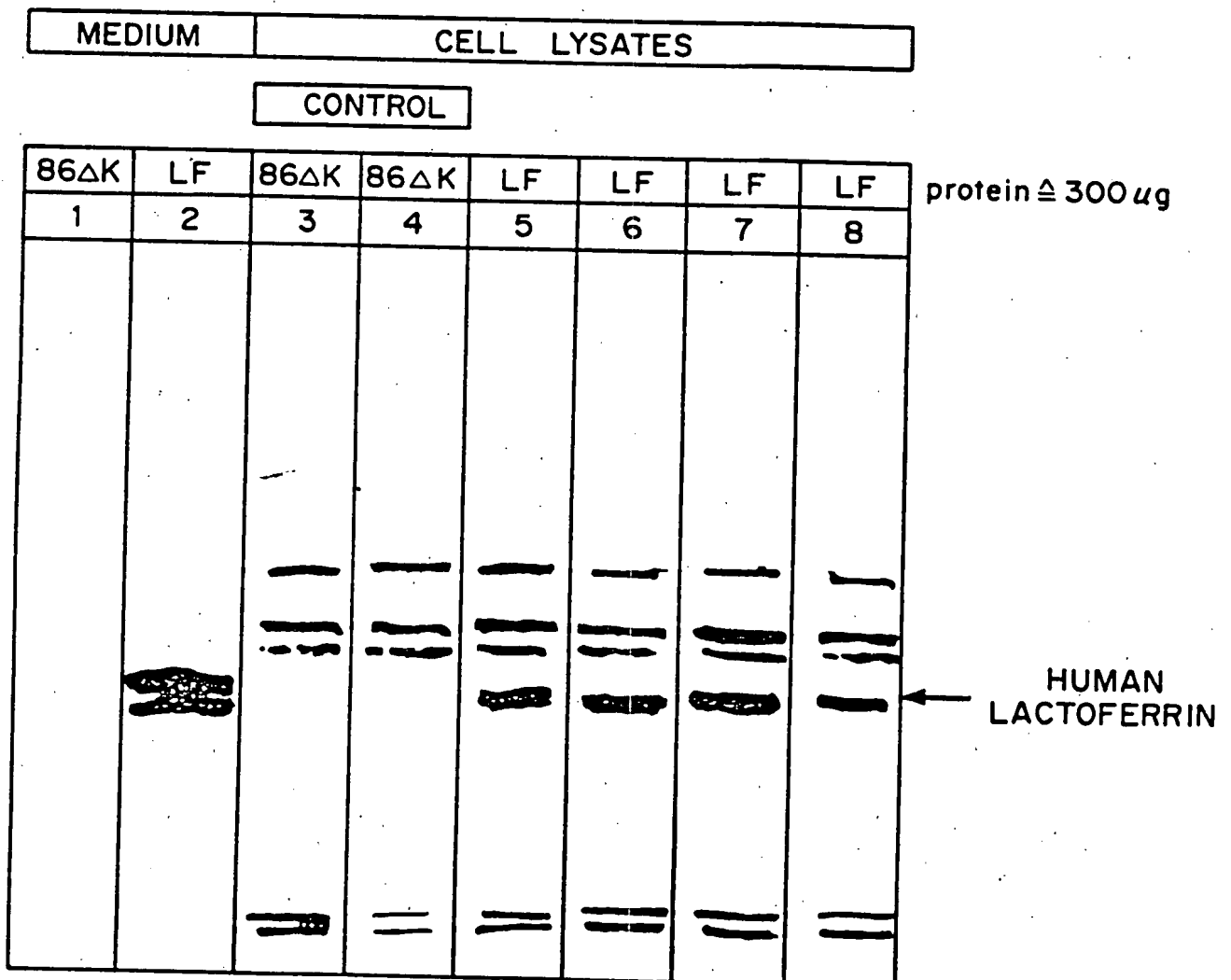
TCCCCTGCTG AAGGTGGGGA TTGCCCATCC ATCTGCTTAC AATTCCCTGC TGTCGTCTTA

2300

GCAAGAAGTA AAATGAGAAA TTTTGTGAA AAAAAAAAAA AAAAAAAAAA AAAAAAAAAA ) 1

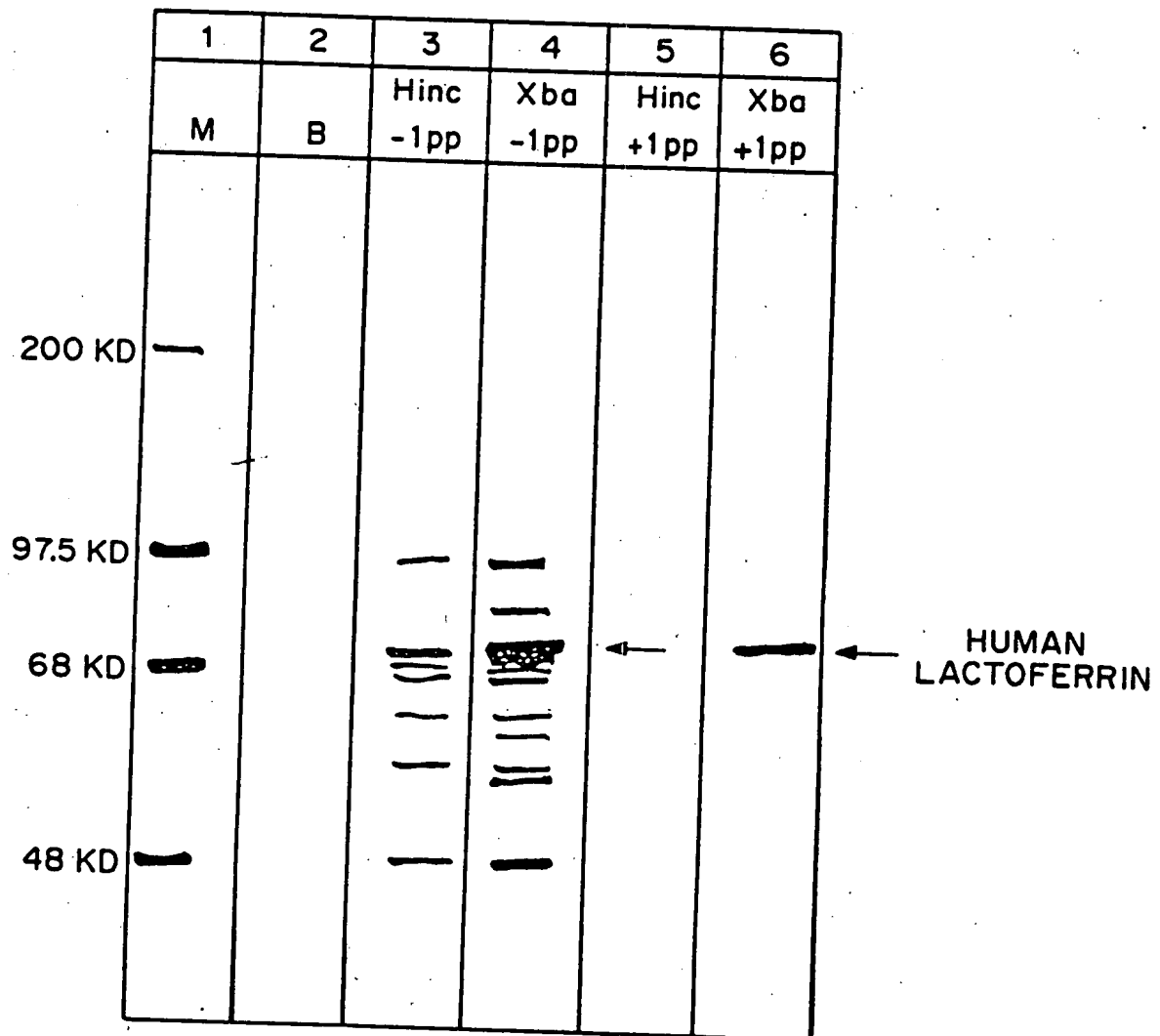
fig. 3

EXPRESSION OF RECOMBINANT HUMAN  
LACTOFERRIN PROTEIN FROM hLF cDNA  
(COS M6 - CELLS)



**fig. 4**

IN VITRO TRANSLATION 2140 bP hLF SEQUENCE



# INTERNATIONAL SEARCH REPORT

International Application No.

PCT/US 90/02356

## I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all) \*

According to International Patent Classification (IPC) or to both National Classification and IPC

IPC<sup>5</sup>: C 12 N 15/12

## II. FIELDS SEARCHED

Minimum Documentation Searched <sup>7</sup>

Classification System

Classification Symbols

IPC<sup>5</sup>

C 12 N, C 12 P

Documentation Searched other than Minimum Documentation  
to the Extent that such Documents are Included in the Fields Searched \*

## III. DOCUMENTS CONSIDERED TO BE RELEVANT \*

Category *	Citation of Document, <sup>11</sup> with indication, where appropriate, of the relevant passages <sup>12</sup>	Relevant to Claim No. <sup>13</sup>
Y	Blood, volume 79, no. 4, 1987, Grune & Stratton, Inc., (New York, US), T.A. Rado et al.: "Isolation of lactoferrin cDNA from a human myeloid library and expression of mRNA during normal and leukemic myelopoiesis", pages 989-993 see the whole article cited in the application	1-6,8,9
Y	European Journal of Biochemistry, volume 145, 1984, FEBS, (Berlin, DE), M.-H. Metz-Boutigue et al.: "Human lactotransferrin: amino acid sequence and structural comparisons with other transferrins", pages 659-676 see the whole article cited in the application	1-6,8,9
X	--	12,13
P,X	American Dairy Science Association and American Society of Animal Science Combined Annual Meeting, ./.	1,2,8,9

\* Special categories of cited documents: <sup>10</sup>

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step

"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"A" document member of the same patent family

## IV. CERTIFICATION

Date of the Actual Completion of the International Search

21st August 1990

Date of Mailing of this International Search Report

04.10.90

International Searching Authority

EUROPEAN PATENT OFFICE

Signature of Authorized Officer

R.J. Eernisse